WEST Search History

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DATE: Wednesday, May 03, 2006

Hide? Set Name Query

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DB=PGPB, USPT; PLUR=YES; OP=OR

☐ L1 (mutant\$ or mutat\$) near2 fie

28

END OF SEARCH HISTORY

First Hit Previous Doc Next Doc Go to Doc#

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L1: Entry 18 of 28 File: PGPB Sep 5, 2002

DOCUMENT-IDENTIFIER: US 20020124282 A1

TITLE: Plant reproduction polynucleotides and methods of use

Summary of Invention Paragraph:

[0004] Arabidopsis fie mutants (for fertilization-independent endosperm) isolated by Ohad et al. (Proc. Natl. Acad. Sci. USA 93:5319-5324, 1996; see also U.S. Pat. No. 6,229,064) exhibit replication of the central cell nucleus, initiating endosperm development, in the absence of fertilization. Inheritance of the mutant fie allele by the female gametophyte results in embryo abortion; thus, the trait can be transmitted to progeny only by the male gametophyte. The Arabidopsis FIE gene was cloned (Ohad et al., The Plant Cell 11:407-416 (1999); GenBank entry AF129516) and found to encode a polypeptide related to the WD Polycomb group proteins encoded by, for example, Esc in Drosophila (Gutjahr et al., EMBO J 14:4296-4306 (1995); Sathe and Harte, Mech. Dev. 52:77-87 (1995); Jones and Gelbart, Mol. Cell. Biol. 13:6357-6366 (1993). WD polycomb proteins may interact with other polynucleotides to form complexes which interfere with gene transcription (Pirrotta, Cell 93:333-336 (1998). Fertilization may trigger alteration of the protein complexes, allowing transcription of genes involved in endosperm development. Thus, loss-of-function fie mutants would lack the ability to form the protein complexes which repress transcription, and endosperm development could proceed independent of fertilization (Ohad et al. 1999, supra).

Detail Description Paragraph:

[0177] Gene inactivation can be used to determine the function of ZmFIE genes in the regulation of endosperm development. When fertilization is prevented in Arabidopsis plants heterozygous for <u>fie mutant</u> alleles, siliques nevertheless elongate and contain seed-like structures due to partial endosperm development. No embryo development is observed (Ohad, Yadegari et al.(1999) Plant Cell 11:407-415). Maize <u>fie mutants</u> would be expected to develop endosperm (or kernels) in the absence of fertilization (i.e. when immature ears are protected from pollination by bags).

Detail Description Paragraph:

[0185] When such nucl:CHD-DR transformation is accomplished in a <u>mutant fie</u> background, both de novo embryo development and endosperm development without fertilization could occur. (see Ohad et al. 1999 The Plant Cell 11:407-415). Upon microscopic examination of the developing embryos it will be apparent that apomixis has occurred by the presence of embryos budding off the nucellus.

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L1: Entry 24 of 28

File: USPT

Dec 7, 2004

DOCUMENT-IDENTIFIER: US 6828477 B1

** See image for Certificate of Correction **

TITLE: Method of enhancing endosperm development in a plant

<u>Detailed Description Text</u> (4):

The present invention is based, at least in part, on the discovery of a set of female-gametophytic mutations, termed fie (fertilization-independent endosperm), and the subsequent cloning of the genes involved. Three mutants are disclosed here fiel, fie2, and fie3, which have been mapped to chromosomes 1, 2, and 3 of Arabidopsis, respectively. The fie mutations affect the central cell, allowing for replication of the central cell nucleus and endosperm development without fertilization. FIE/fie seed coat and fruit undergo fertilization-independent differentiation, showing that the fie female gametophyte is the source of signals that activates sporophytic fruit and seed coat development. Generally, the mutant fie alleles are not transmitted by the female gametophyte. Inheritance of a mutant fie allele (e.g., fie3) by the female gametophyte usually results in embryo abortion, even when the pollen bears the wild-type FIE allele. In the case of fiel and fie2, however, transmission of the trait occurs in about 1% of the progeny from the female gametophyte. In contrast, the fiel, fie2, and fie3 mutant alleles are passed through the male gametophyte (i.e., pollen) in normal fashion.

Detailed Description Text (23):

Gene expression can be inactivated using recombinant DNA techniques by transforming plant cells with constructs comprising transposons or T-DNA sequences. <u>FIE mutants</u> prepared by these methods are identified according to standard techniques. For instance, mutants can be detected by PCR or by detecting the presence or absence of FIE mRNA, e.g., by Northern blots. Mutants can also be selected by assaying for development of endosperm in the absence of fertilization.

Detailed Description Text (37):

As noted above, FIE proteins as products of polycomb group genes are believed to form large complexes in vivo. Thus, production of dominant-negative forms of FIE polypeptides that are defective in their abilities to bind to other polycomb group proteins is a convenient means to inhibit endogenous FIE activity. This approach involves transformation of plants with constructs encoding mutant FIE polypeptides that form defective complexes with endogenous polycomb group proteins and thereby prevent the complex from forming properly. The mutant polypeptide may vary from the naturally occurring sequence at the primary structure level by amino acid substitutions, additions, deletions, and the like. These modifications can be used in a number of combinations to produce the final modified protein chain. Use of dominant negative mutants to inactivate target genes is described in Mizukami et al. Plant Cell 8:831-845 (1996).

Detailed Description Text (61):

The following example describes methods used to identify the <u>fie mutants</u>. The methods described here are generally as described in Ohad et al., Proc. Natl. Acad. Sci. USA 93:5319-5324 (1996).

Detailed Description Text (67):

Heterozygous FIE/fie (Landsberg erecta ecotype) plants were crossed as males with

female plants (Columbia ecotype). Because the <u>mutant fie</u> allele is only transmitted through the male gametophyte, FIE/fie progeny were crossed as males a second time to female gl1/gl1 (Columbia ecotype) plants. Approximately fifty-five progeny were scored for the segregation of the wild-type <u>FIE and mutant fie</u> alleles and for alleles of molecular markers as described previously (Bell, C., et al., Genomics 19: 137-144 (1994)). This analysis indicated that fie3 is located at approximately position 30 on chromosome three, fie2 is located at approximately position 65 on chromosome two, and fie1 is located at approximately position 2 on chromosome one. Genetic recombination frequencies and map distances were calculated according to Koornneef and Stam (Koornneef, M., et al., Methods in Arabidopsis Research, pp. 83-99 (1992)) and Kosambi (Kosambi, Ann. Eugen., 12: 172-175 (1944)).

Detailed Description Text (78):

Seed Coat and Silique Development In a representative line chosen for further study, heterozygous plants produced by back crosses to wild-type plants generated elongated siliques after anther removal with numerous seed-like structures. These results indicated that heterozygous mutant plants were capable of silique elongation and seed-like structure development in the absence of fertilization. We compared the development of the mutant seed-like structures to that of wild-type seeds. After fertilization, the endosperm nucleus replicated and daughter nuclei migrated into the expanding central cell. Ultimately, a syncytium of endosperm nuclei was produced. Nuclear divisions of the endosperm preceded the zygotic divisions that formed the globular stage embryo. Embryo, endosperm or seed coat development did not occur in wild-type plants in the absence of fertilization. Development of the ovule and female gametophyte in heterozygous mutant plants was normal. Just prior to flower opening, female gametophytes in these plants contained a single, prominent central cell nucleus. Subsequently, in the absence of fertilization, central cells with two large nuclei were detected. Further divisions resulted in the production of additional nuclei that migrated into the expanded central cell. Later in development, a nuclear-syncytium was formed with abundant endosperm nuclei. These results indicated that the central cell in mutant female gametophytes initiated endosperm development in the absence of fertilization. We have named this mutation fie for fertilization-independent endosperm. By contrast, replication of other nuclei in fie female gametophytes (egg, synergid, or antipodal) was not detected. Thus, the fie mutation specifically affects replication of the central cell nucleus.

Detailed Description Text (79):

We analyzed the frequency of multinucleate central cell formation in fie female gametophytes by comparing the percentage of multinucleate central cells at three, five, and six days after emasculation of heterozygous FIE/fie and control wild-type flowers. At each time point, only 3% to 5% of wild-type central cells had more than one nucleus. Because none had more than two nuclei, most likely, these represented central cells with haploid nuclei that had not fused during female gametophyte development. By contrast, the percentage of central cells in female gametophytes from FIE/fie siliques with two or more nuclei increased from 21% to 47% over the same time period. These results indicated that the fie mutation caused a significant increase in formation of multinucleate central cells in the absence of fertilization. The fact that close to 50% of the female gametophytes in heterozygous plants had multinucleate central cells suggested that fie is a gametophytic mutation because a 1:1 segregation of wild-type and mutant fie alleles occurs during meiosis.

Detailed Description Text (80):

We compared the fertilization-independent development of the maternal seed coat in FIE/fie seed-like structures to that of fertilized wild-type seeds. The seed coat in wild-type Arabidopsis is generated by the integuments of the ovule and surrounds the developing embryo and endosperm. Similarly, FIE/fie ovule integuments formed a seed coat that surrounded the developing mutant endosperm. These results indicated that the <u>fie mutation</u> activated both endosperm development and maternal sporophytic

seed coat and silique differentiation that support reproduction. No other effects on sporophytic growth and development were detected in FIE/fie plants.

Detailed Description Text (82):

To understand the mode of inheritance of the fie mutation, we analyzed the progeny of reciprocal crosses. FIE3/fie3 females, crossed to wild-type males, produced siliques with approximately equal numbers of viable seeds with normal green embryos and nonviable white seeds with embryos aborted at the heart stage (344:375, 1:1, c2=1.3, P>0.2). Viable seeds from this cross were germinated and all 120 F1 progeny generated were wild-type. That is, none of the F1 progeny had significant levels of F2 aborted seeds in their siliques after self-pollination. Nor did the F1 progeny demonstrate fertilization-independent development. This indicated that presence of the fie mutant allele in the female gametophyte, even when the male provided a wild-type allele, resulted in embryo abortion. Thus, the fie mutation is not transmitted by the female gametophyte to the next generation. To study transmission of fie through the male gametophyte, we pollinated female wild-type plants with pollen from male FIE3/fie3 plants. Siliques from these crosses contained no aborted F1 seed. F1 plants were examined and a 1:1 segregation of wild-type and FIE3/fie3 genotype was observed (62:58, c2=0.13, P>0.5). This indicated that wild-type and mutant fie3 alleles were transmitted by the male gametophyte with equal efficiency. That is, fie does not affect male gametophyte, or pollen grain, function. Results from reciprocal crosses were verified by analyzing the progeny from self-pollinated FIE3/fie3 plants. Self-pollinated siliques displayed 1:1 segregation of normal and aborted seeds (282:286, c2=0.03, P>0.8). Viable seed from self-pollinated siliques were germinated and a 1:1 (71:64, c2 0.36, P>0.5) segregation of wild-type and FIE3/fie3 progeny was observed. These results confirmed that inheritance of a fie mutant allele by the female gametophyte resulted in embryo abortion, and that inheritance of a fie mutant allele by the male gametophyte did not affect pollen function. Thus, the wild-type FIE3 allele probably carries out a function unique to the female gametophyte and does not appear to be needed for male fertility.

Detailed Description Text (85):

In wild-type plants, fertilization initiates embryogenesis and endosperm formation, and activates maternal seed coat and silique development. The results presented here indicate that specific aspects of plant reproductive development can occur in FIE/fie plants in the absence of fertilization. These include silique elongation, seed coat formation, and endosperm development. Morphological analysis shows that early aspects of fertilization-independent fie endosperm development closely resemble fertilized wild-type endosperm development. First, the fie central cell nucleus is stimulated to undergo replication. Second, nuclei that are produced migrate from the micropylar end of the central cell and take up new positions in the central cell. Third, the developing fie central cell expands to form an endosperm cavity. Thus, the requirement for fertilization to initiate these early events in endosperm formation has been eliminated by the fie mutation. This suggests that FIE plays a role in a signal transduction pathway that links fertilization with the onset of central cell nuclear replication and early endosperm development.

Detailed Description Text (87):

One can envision two possible mechanisms for how FIE regulates replication of the central cell nucleus in response to fertilization. The protein encoded by the FIE gene may be involved in a positive regulatory interaction. In this model, FIE is required for the central cell to initiate endosperm development. Normally, fertilization is needed for the presence of active FIE protein. The <u>fie mutation</u> results in the presence of active protein in the absence of fertilization. Alternatively, F1 may by involved in a negative regulatory interaction. In this model, the function of FIE protein is to prevent the central cell from initiating endosperm development, and fertilization results in the inactivation of FIE protein. The <u>fie mutation</u> results in the production of inactive protein, so that fertilization is no longer required to initiate endosperm development However,

complementation experiments using transgenic plants indicate that FIE1 and FIE3 alleles are dominant over their respective mutant alleles. This indicates that the wild-type allele is involved in a negative regulatory interaction. Recently, it has been shown that cyclin-dependent kinase complexes, related to those that function in mammals, control the induction of DNA synthesis and mitosis in maize endosperm (Grafi, G. et al., Science 269: 1262-1264 (1995)). Because fie stimulates replication of the central cell, fie may, either directly or indirectly, impinge upon cell cycle control of the central cell nucleus, allowing replication to take place in the absence of fertilization.

Detailed Description Text (89):

The analysis of FIE/fie mutant plants has provided clues about interactions between endosperm and maternal sporophytic tissues. FIE/fie ovule intequments surrounding a mutant fie female gametophyte initiate seed coat development, whereas FIE/fie integuments in contact with a quiescent wild-type female gametophyte do not develop. This suggests that the FIE/fie ovule integuments initiate seed coat differentiation in response to a signal produced by the fie female gametophyte. We propose that the source of the signal is the mutant fie central cell that has initiated endosperm development, although we cannot rule out the participation of other cells in the fie female gametophyte. In wild-type plants, most likely, fertilization of the central cell produces an endosperm that activates seed coat development. This is consistent with experiments showing that the maize endosperm interacts with nearby maternal cells (Miller, M. E., et al., Plant Cell 4: 297-305 (1992)). FIE/fie plants also display fertilization-independent elongation of the ovary to form the silique. We propose that a signal is produced by the developing seed-like structures to initiate silique elongation. This is in agreement with experiments suggesting that seeds are the source of hormones, auxins and gibberellins, that activate fruit development (Lee, T. D. Plant Reproductive Ecology, pp. 179-202 (1988)). Taken together, these results suggest that the fertilized female gametophyte activates maternal developmental programs.

Detailed Description Text (91):

Certain plant species display aspects of fertilization-independent reproductive development, including apomictic generation of embryo and endosperm, and development of the maternal seed coat and fruit (reviewed in (Koltunow, a. Plant Cell 5: 1425-1437 (1993)). The <u>fie mutation</u> reveals that Arabidopsis, a sexually reproducing plant, has the genetic potential for aspects of fertilization-independent reproductive development. It is not known whether the mechanism of fertilization-independent endosperm development conferred by the <u>fie mutation</u> is the same as autonomous endosperm formation observed in certain apomictic plant species. However, the fact that the fie phenotype is caused by a single genetic locus substantiates the view that the number of genetic differences between sexually and asexually reproducing plants is small (Koltunow, a M., et al., Plant Physiol 108:1345-1352 (1995)).

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    Human SRCAP and Drosophila melanogaster DOM are homologs that function in
ΤI
    the Notch signaling pathway
    Eissenberg, Joel C.; Wong, Madeline; Chrivia, John C.
ΑU
    Edward A. Doisy Department of Biochemistry and Molecular Biology, Saint
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    Louis University School of Medicine, St. Louis, MO, 63104, USA
    Molecular and Cellular Biology (2005), 25(15), 6559-6569
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    Evaluation of ***female***
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CS
     Khon Kaen Univ, Fac Med, Dept Obstet and Gynecol, Khon Kaen 40002,
     Thailand
    yutwer@kku.ac.th
     Journal of the Medical Association of Thailand, (AUG 2005) Vol. 88, No. 8,
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     CODEN: JMTHBU. ISSN: 0125-2208.
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    2002:392271 CAPLUS
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    Protein-binding RNA sequences for incorporation into into mRNAs and their
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     Conner, Timothy W.; Fabbri, Bradon J.; Huang, Jintai
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     U.S. Pat. Appl. Publ., 69 pp.
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CODEN: USXXCO

DT Patent

LA English

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L7 ANSWER 6 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2002:937303 CAPLUS

DN 138:20443

TI Endocrine disruptor screening using DNA chips of endocrine disruptor-responsive genes

L7 ANSWER 7 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2002:560356 CAPLUS

DN 137:245889

TI Pituitary hypoplasia and lactotroph dysfunction in mice deficient for cyclin-dependent kinase-4

L7 ANSWER 8 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1

AN 2002:916162 CAPLUS

DN 138:201982

TI The E2F cell cycle regulator is required for Drosophila nurse cell DNA replication and apoptosis

L7 ANSWER 9 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2002:833756 CAPLUS

DN 138:266045

TI Prospects for using genetic transformation for improved SIT and new biocontrol methods

AU Handler, Alfred M.

CS Center for Medical, Agricultural, and Veterinary Entomology, Agricultural Research Service, US Department of Agriculture, Gainesville, FL, 32608, USA

SO Genetica (Dordrecht, Netherlands) (2002), 116(1), 137-149 CODEN: GENEA3; ISSN: 0016-6707

L7 ANSWER 10 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2000:690493 CAPLUS

DN 133:361422

TI Activation-induced nuclear translocation of RING3

AU Guo, Ning; Faller, Douglas V.; Denis, Gerald V.

CS Cancer Research Center, Boston University School of Medicine, Boston, MA, USA

SO Journal of Cell Science (2000), 113(17), 3085-3091 CODEN: JNCSAI; ISSN: 0021-9533

L7 ANSWER 11 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 2

AN 1996:381446 CAPLUS

DN 125:106425

TI The DNA-binding and enhancer-blocking domains of the Drosophila suppressor of hairy-wing protein

L7 ANSWER 12 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 3

AN 1996:722419 CAPLUS

DN 126:87263

TI Myc and Max homologs in Drosophila

L7 ANSWER 13 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 4

AN 1995:547407 CAPLUS

DN 123:103538

TI Structure and expression of the br-c locus in Drosophila melanogaster (Diptera: Drosophilidae)

L7 ANSWER 14 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1993:666326 CAPLUS

DN 119:266326

TI Binary microbe system for biological control of Fusarium wilt of tomato. Enhanced root-colonization of an antifungal rhizoplane bacterium supported by a chitin-degrading bacterium

- AU Toyoda, Hideyoshi; Morimoto, Masayuki; Kakutani, Koji; Morikawa, Masaaki; Fukamizo, Tamo; Goto, Sachio; Terada, Hikojiro; Ouchi, Seiji
- CS Fac. Agric., Kinki Univ., Nara, 631, Japan
- SO Nippon Shokubutsu Byori Gakkaiho (1993), 59(4), 375-86 CODEN: NSBGAM; ISSN: 0031-9473
- L7 ANSWER 15 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN
- AN 1993:402480 CAPLUS
- DN 119:2480
- TI Trans-splicing ribozymes, their preparation, and their use in cell ablation
- IN Haseloff, James; Brand, Andrea; Perrimon, Norbert; Goodman, Howard M.
- PA General Hospital Corp., USA; Harvard College
- SO PCT Int. Appl., 112 pp.

CODEN: PIXXD2

DT Patent

LA English

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- L7 ANSWER 16 OF 22 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
- AN 1991:186709 BIOSIS
- DN PREV199191101458; BA91:101458
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- AN 1982:153698 BIOSIS
- DN PREV198273013682; BA73:13682
- TI THE TITANIUM SILICONE RUBBER CLIP FOR ***FEMALE***

 STERILIZATION .
- L7 ANSWER 20 OF 22 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
- AN 1982:175712 BIOSIS
- DN PREV198273035696; BA73:35696
- TI LUTEAL PHASE PREGNANCIES IN ***FEMALE*** ***STERILIZATION***
 PATIENTS.
- L7 ANSWER 21 OF 22 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN

- AN 1981:164577 BIOSIS
- DN PREV198171034569; BA71:34569
- TI TECHNICAL FAILURES IN TUBAL RING STERILIZATION INCIDENCE PERCEIVED REASONS OUTCOME AND RISK FACTORS.
- L7 ANSWER 22 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN
- AN 1965:76327 CAPLUS
- DN 62:76327
- OREF 62:13549f-q
- TI Evidence for common control of tyrosinase and L-amino acid oxidase in Neurospora
- AU Horowitz, N. H.
- CS California Inst. of Technol., Pasadena
- SO Biochemical and Biophysical Research Communications (1965), 18(5-6),

686-92

AΒ

CODEN: BBRCA9; ISSN: 0006-291X

=> d 17 ab 5 9 10 14 15 22

L7 ANSWER 5 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN

The methods and materials disclosed herein are directed to the control of AB gene expression in plants by means of translational repression. RNA-binding proteins binding specifically to ***operator*** sequences positioned in the 5' untranslated region of an MRNA reduce translation. Such translation repression systems are useful, for example, for reducing expression of an herbicide-tolerance gene in reproductive tissues of a plant that retains vegetative tolerance. Application of the herbicide renders the plant male- or ***female*** - ***sterile*** . The use the MS2 and Q.beta. coat proteins and Saccharomyces cerevisiae ribosomal protein L32 as RNA binding proteins capable of inhibiting translation is demonstrated in vitro. Translation of .beta.-glucuronidase mRNA carrying the MS2 translational ***operator*** in a wheat germ system was effectively inhibited by addn. of mRNA for the coat protein. The protein was also an effective inhibitor in corn, wheat and tobacco leaf protoplasts. Expression of the genes for MS2 coat protein from pollen-specific promoters is demonstrated in transgenic corn. These plants also carried the gene for a glyphosate-resistant EPSP synthase contg. the MS2 coat protein-binding translation repressor. Pollen were sensitive to glyphosate whereas the plant was resistant, meaning that male sterility could be induced by treatment with the herbicide.

L7 ANSWER 15 OF 22 CAPLUS COPYRIGHT 2006 ACS on STN

A chimeric trans-splicing ribozyme comprising a first RNA sequence functioning in targeting of the ribozyme, and a 2nd RNA sequence which is capable of being transferred into the target RNA as a result of the ribozyme activity is described. The first RNA sequence hybridizes to an RNA encoding a ***transcription*** ***activator*** Expression of the RNA or DNA encoding this ribozyme is operably linked to expression of the ***transcription*** ***activator*** protein. Expression of the ribozyme-encoding gene in a multicellular organism provides a means of specific cell ablation, and can be used to produce ***female*** ***sterility*** in plants, or to immunize plants against a pathogen. The ribozyme can be expressed as a proribozyme, which contain self-complementary sequences which prevent self-cleavage. In the presence of target sequences, the intramol. interaction is inhibited due to preferential interaction with the substrate nucleic acid and the proribozyme is activated. Trans-splicing ribozymes, directed to coat protein RNA of cucumber mosaic virus, and capable of inserting RNA encoding the A chain of diphtheria toxin into the coat protein RNA, were described. A ribozyme capable of inserting the toxin mRNA into the GAL4 mRNA was prepd. and expressed in a cell-specific manner in Drosophila in order to study the effects of cell-specific ablation on development.

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PROCESSING COMPLETED FOR L8

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L9
=> s 19 and arabidopsis
L10
            10 L9 AND ARABIDOPSIS
=> d l10 1-10
     ANSWER 1 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
L10
AN
     2004:691479 CAPLUS
DN
     141:186046
TI
     Use of
              ***Arabidopsis***
                                  thaliana leafy cotyledon 1 gene for
     modulating embryo development in transgenic plants
IN
     Robert L.; Bui, Anhthu; Kwong, Raymond
PΑ
SO
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Harada, John; Lotan, Tamar; Ohto, Masa-Aki; Goldberg, Robert B.; Fischer,

The Regents of the University of California, USA

U.S., 38 pp., Cont.-in-part of U.S. 6,320,102.

CODEN: USXXAM

DТ Patent

LA English

FAN.CNT 7

	PA	TENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	יייי	6781035	B1	20040824	US 2000-516052	20000301
r T		6545201	B1	20030408	US 1998-26221	19980219
	US	6235975	B1	20010522	US 1998-103478	19980624
	US	6320102	B1	20011120	US 1998-193931	19981117
	CA	2399886	AA	20010907	CA 2001-2399886	20010221
	WO	2001064022	A2	20010907	WO 2001-US5454	20010221
	AU 2001041600		A5 200	10912 AU 2001-41600	20010221	
	ΕP	1263280	A1	20021211	EP 2001-912861	20010221
	JP 2004500823		T2 2004	0115 JP 2001-562933	20010221	
PRAI	US	1997-804534	A2	19970221		
	US	1998-26221	A2	19980219		
	US	1998-103478	A2	19980624		
	US	1998-193931	A2	19981117		
	US	2000-516052	Α	20000301		
	WO	2001-US5454	W	20010221		

L10 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:270094 CAPLUS

DN 140:265609

TI Methods for producing infertile transgenic seeds

IN Mascia, Peter

PΑ Ceres, Inc., USA

SO PCT Int. Appl., 42 pp.

CODEN: PIXXD2

DTPatent

English LΑ

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	WO 2004027038	A2	20040401	WO 2003-US29691	20030917

L10 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:261449 CAPLUS

DN 140:401225

Isolation of the gene encoding Carrot leafy cotyledon1 and expression TIanalysis during somatic and zygotic embryogenesis

ΑU Yazawa, Katsumi; Takahata, Kiminori; Kamada, Hiroshi

Institute of Biological Sciences, Gene Research Center, University of CS Tsukuba, Tsukuba, Ibaraki, 305-8572, Japan

SO Plant Physiology and Biochemistry (Amsterdam, Netherlands) (2004), 42(3), 215-223

CODEN: PPBIEX; ISSN: 0981-9428

L10 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN

AN2003:276758 CAPLUS

DN 138:282470

TI Leafy cotyledon1 gene of ***Arabidopsis*** thaliana and its use in modulating gene expression during embryo development

IN Harada, John J.; Lotan, Tamar; Ohto, Masa-aki; Goldberg, Robert B.;

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Fischer, Robert L.
PA
     The Regents of the University of California, USA
SO
     U.S., 22 pp., Cont.-in-part of U.S. Ser. No. 804,534.
    CODEN: USXXAM
DT
     Patent
LA
    English
FAN.CNT 7
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    US 6545201
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EP 1998-907487
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    EP 977836
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                        A2
     US 1998-193931
                              19981117
L10
    ANSWER 5 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
AN
     2001:843888 CAPLUS
DN
    135:369339
    Leafy cotyledon1 gene of ***Arabidopsis***
ΤI
                                                thaliana and their uses in
    modulating gene expression during embryo development
IN
    Harada, John J.; Lotan, Tamar; Ohto, Masa-aki; Goldberg, Robert B.;
    Fischer, Robert L.
PΔ
    Regents of the University of California, USA
SO
    U.S., 31 pp., Cont.-in-part of U.S. 6,235,975.
    CODEN: USXXAM
DT
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    English
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    US 6320102
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                                        US 1998-193931
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    US 6545201
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                           20030408
                                       US 1998-26221
                                                              19980219
                B1 20010522 US 1998-103478
A2 19991229 WO 1999-US14384
    US 6235975
                                         US 1998-103478 19980624
WO 1999-US14384 19990624
    WO 9967405
            AU 9948313
                              A1 20000110
                                               AU 1999-48313
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                                        US 2000-516052
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    US 1998-103478
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    US 1998-193931
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                              19981117
    WO 1999-US14384
                        W
                              19990624
    ANSWER 6 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
L10
    2001:661191 CAPLUS
AN
    135:222367
DN
ΤI
    Leafy cotyledon1 genes and promoter from
                                             ***Arabidopsis***
                                                               thaliana
    and their uses in embryogenesis in plants
IN
    Harada, John; Lotan, Tamar; Ohto, Masa-aki; Goldberg, Robert B.; Fischer,
    Robert L.; Bui, Anhthu; Kwong, Raymond
PΑ
    Regents of the University of California, USA
SO
    PCT Int. Appl., 73 pp.
    CODEN: PIXXD2
DT
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    English
LΑ
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    PATENT NO.
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    WO 2001064022
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PΙ
                                         WO 2001-US5454
                             20010907
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                                                              20010221
                              B1 20040824 US 2000-516052
           US 6781035
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    CA 2399886
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                0 A5
A1
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    US 1998-26221
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A2
      US 1998-103478
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      US 1998-193931
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                                      19981117
      WO 2001-US5454
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      ANSWER 7 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
      2001:366745 CAPLUS
      134:362280
        ***Arabidopsis*** thaliana gene LEAFY-COTYLEDON1, its DNA and cDNA
      sequences, promoter and use in modulating embryo development in transgenic
      plants
      Harada, John J.; Lotan, Tamar; Ohto, Masa-aki; Goldberg, Robert B.;
      Fischer, Robert L.
      Regents of the University of California, USA
      U.S., 32 pp., Cont.-in-part of U.S. Ser. No. 26,221.
      CODEN: USXXAM
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FAN.CNT 7
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PI US 6235975 B1 20010522 US 1998-103478 19980624
    US 6545201 B1 20030408 US 1998-26221 19980219
    US 6320102 B1 20011120 US 1998-193931 19981117
    WO 9967405 A2 19991229 WO 1999-US14384 19990624
    AU 9948313 A1 20000110 AU 1999-48313 1
US 6781035 B1 20040824 US 2000-516052 20000301
PRAI US 1997-804534 B2 19970221
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    US 1998-193931 A 19981117
    WO 1999-US14384 W 19990624
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                                                                                          19990624
     ANSWER 8 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
      2000:404694 CAPLUS
      133:132589
      Changes in gene expression in the leafy cotyledon1 (lec1) and fusca3
      (fus3) mutants of ***Arabidopsis*** thaliana L.
      Vicient, Carlos M.; Bies-Etheve, Natacha; Delseny, Michel
      Laboratoire de Physiologie et Biologie Moleculaire des Plantes, Centre
      National de la Recherche Scientifique UMR 5545, Universite de Perpignan,
      Perpignan, 66860, Fr.
      Journal of Experimental Botany (2000), 51(347), 995-1003
      CODEN: JEBOA6; ISSN: 0022-0957
L10 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
      1999:819513 CAPLUS
      132:60145
      Leafy cotyledon1 gene and promoter from ***Arabidopsis*** thaliana and
      their uses for embryo-specific gene expression
      Harada, John J.; Lotan, Tamar; Ohto, Masa-aki; Goldberg, Robert B.;
      Fischer, Robert L.
      Regents of the University of California, USA
      PCT Int. Appl., 69 pp.
      CODEN: PIXXD2
      Patent
      English
FAN.CNT 7
                        KIND DATE
                                                APPLICATION NO.
      PATENT NO.
                                                                              DATE
                             A2 19991229 WO 1999-US14384 19990624
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      WO 9967405
              ~US_6235975
                                       B1 20010522 US 1998-103478
                                                                                           19980624
      US 6320102 B1 20010522 US 1998-103478

US 6320102 B1 20011120 US 1998-193931 19981117

AU 9948313 A1 20000110 AU 1999-48313 19990624

US 1998-103478 A 19980624

US 1998-193931 A 19981117

US 1997-804534 B2 19970221

US 1998-26221 A2 19980219
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PRAI US 1998-103478
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      WO 1999-US14384
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L10 ANSWER 10 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
      1998:604991 CAPLUS
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129:198906

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TI
     Leafy cotyledon1 gene and promoter from ***Arabidopsis*** thaliana and
     their uses for embryo-specific gene expression
IN
     Harada, John J.; Lotan, Tamar; Ohto, Masa-aki; Goldberg, Robert B.;
     Fischer, Robert L.
PA
     The Regents of the University of California, USA
SO
     PCT Int. Appl., 55 pp.
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    PATENT NO.
    WO 9837184 A1
    WO 9837184 A1 19980827 WO 1998-US2998 19980220
CA 2281487 AA 19980827 CA 1998-2281487
AU 9863283 A1 19980909 AU 1998-63283 19980220
AU 732026 B2 20010412
EP 977836 A1 20000209 EP 1998-907487 19980220
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    PRAI US 1997-804534 A 19970221
US 1998-26221 A 19980219
WO 1998-US2998 W 19980220
=> d l10 8 ab
=> s lec1/ab, bi and 12
      2 LEC1/AB, BI AND L2
L11
=> d l11 1-2
L11 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
AN
    2006:332217 CAPLUS
DN
    144:325851
ΤI
    Use of controlled induction of ***seed***
                                                   ***sterility***
    controlling the spread of transgenic plants
IN
    Mascia, Peter N.
PA
    Ceres, Inc., USA
so
    PCT Int. Appl., 43 pp.
    CODEN: PIXXD2
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LA
    English
FAN.CNT 1
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ΡI
    WO 2006009922
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                                                                20050620
L11 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
    2004:270094 CAPLUS
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    140:265609
TI Methods for producing infertile transgenic seeds
IN
    Mascia, Peter
PA
    Ceres, Inc., USA
SO
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    WO 2004027038
PΙ
                        A2 20040401 WO 2003-US29691
                                                                  20030917
    WO 2004027038
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           20 ((MUTANT? OR MUTAT?)(5A) FIE)/AB, BI
=> s 112 and 12
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=> s l13 not l11
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L15
     2001:8671 CAPLUS
AN
DN
     134:190759
ΤI
    Hypomethylation promotes autonomous endosperm development and rescues
     postfertilization lethality in
                                     ***fie***
                                                   ***mutants***
     Vinkenoog, Rinke; Spielman, Melissa; Adams, Sally; Fischer, Robert L.;
ΑU
     Dickinson, Hugh G.; Scott, Rod J.
    Department of Biology and Biochemistry, University of Bath, Bath, BA2 7AY,
CS
so
     Plant Cell (2000), 12(11), 2271-2282
     CODEN: PLCEEW; ISSN: 1040-4651
    ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 2
L15
     1996:326996 CAPLUS
AN
TΙ
    A mutation that allows endosperm development without fertilization
    Ohad, Nir; Margossian, Linda; Hsu, Yung-Chao; Williams, Chad; Repetti,
ΑU
     Peter; Fischer, Robert L.
CS
    Dep. Plant Biology, Univ. California, Berkeley, CA, 94720-3102, USA
     Proceedings of the National Academy of Sciences of the United States of
SO
                                                                          printed
    America (1996), 93(11), 5319-5324
     CODEN: PNASA6; ISSN: 0027-8424
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     1996:136937 CAPLUS
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                                                                                   prated
DN
     124:280540
TI
    AINTEGUMENTA, an APETALA2-like gene of Arabidopsis with pleiotropic roles
     in ovule development and floral organ growth
ΑU
     Elliott, Robert C.; Betzner, Andreas S.; Huttner, Eric; Oakes, Marie P.;
     Tucker, William Q. J.; Gerentes, Denise; Perez, Pascual; Smyth, David R.
CS
    Dep. Genet. Dev. Biol., Monash Univ., Clayton, 3168, Australia
     Plant Cell (1996), 8(2), 155-68
     CODEN: PLCEEW; ISSN: 1040-4651
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STN INTERNATIONAL LOGOFF AT 14:53:27 ON 03 MAY 2006